

**Claim Amendments**

1. **(previously presented)** A pharmaceutical composition for intramammary administration to a non-human mammal, wherein:  
the composition comprises:  
an antibacterial agent,  
prednisolone, and  
a pharmaceutically acceptable carrier; and  
the composition comprises at least 20 mg of prednisolone per unit dose.
2. **(previously presented)** The composition according to claim 1, wherein the composition comprises prednisolone in an amount of 20 to 40 mg per unit dose.
3. **(previously presented)** The composition according to claim 2, wherein the composition comprises prednisolone in an amount of 20 to 30 mg per unit dose.
4. **(previously presented)** The composition according to claim 1, wherein the antibacterial agent is a cephalosporin.
5. **(previously presented)** The composition according to claim 4, wherein the cephalosporin is cephapirin.
6. **(previously presented)** The composition according to claim 4, wherein the cephalosporin is cefquinome.
7. **(previously presented)** The composition according to claim 1, wherein the composition comprises the antibacterial agent in an amount of 10 to 500 mg per unit dose.
8. **(withdrawn)** A process for preparing a pharmaceutical composition according to claim 1, comprising the steps of mixing an oil and one or more pharmaceutically acceptable

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additives to form a carrier, and suspending the antibacterial agent and the prednisolone in the carrier.

**9. (Canceled).**